

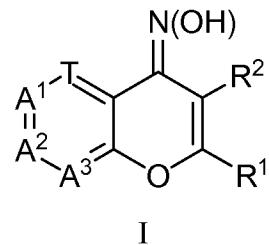
Applicants: Jeremy Green et al.
Application No.: 10/808,678

THE CLAIMS

Please replace all prior versions and listings of claims with the amended claims as follows:

1-46. (Canceled)

47. (Currently amended) A pharmaceutical composition comprising a compound of formula I:



I

or a pharmaceutically acceptable salt thereof, and a pharmaceutically acceptable carrier, adjuvant, or vehicle, wherein:

R¹ is Ar¹;

R² is hydrogen;

T is CH;

A¹ is C-halogen, C-CN, or C-R;

each of A² and A³ is, independently, CR⁴;

R⁴ is selected from halogen, NO₂, CN, -(L)_mR, -(L)_mAr¹, or -(L)_mCy¹; or

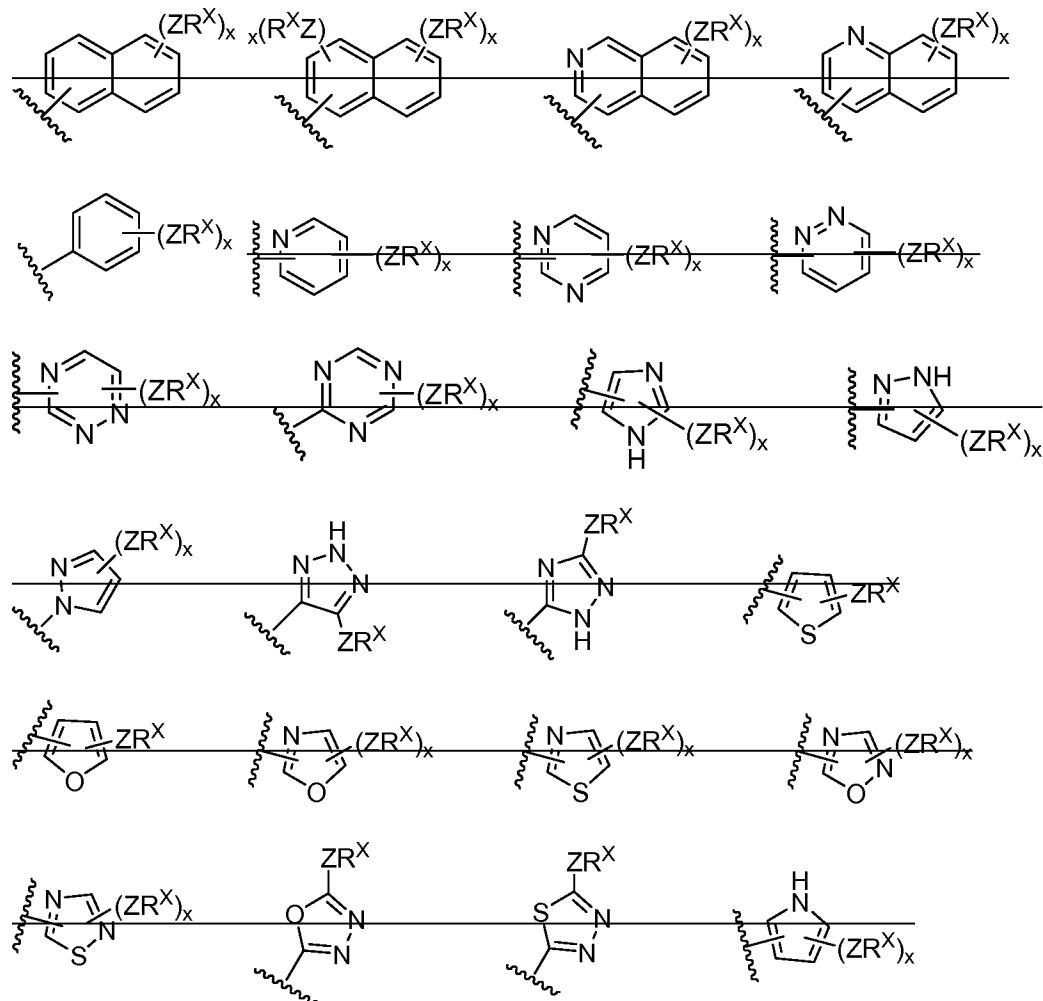
two R⁴ groups on adjacent atoms are taken together to form an optionally substituted 5-7 membered partially unsaturated or fully unsaturated ring having 0-3 heteroatoms independently selected from oxygen, sulfur, or nitrogen, wherein[[:]] each ring formed by two R⁴ groups on adjacent atoms taken together is optionally substituted with up to 4 occurrences of Z-R^X;

Applicants: Jeremy Green et al.
Application No.: 10/808,678

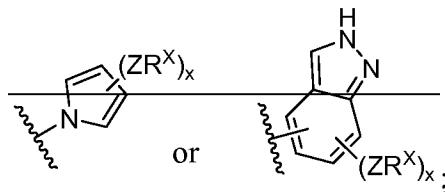
L is a C₁₋₆ alkylidene chain wherein one ~~up to two non adjacent~~ methylene unit of L is optionally replaced by -O-, -N(R)-, -N(R)C(O)-, -C(O)-, -C(O)N(R)-, -SO₂N(R)-, or -N(R)SO₂-;

m is 0 or 1;

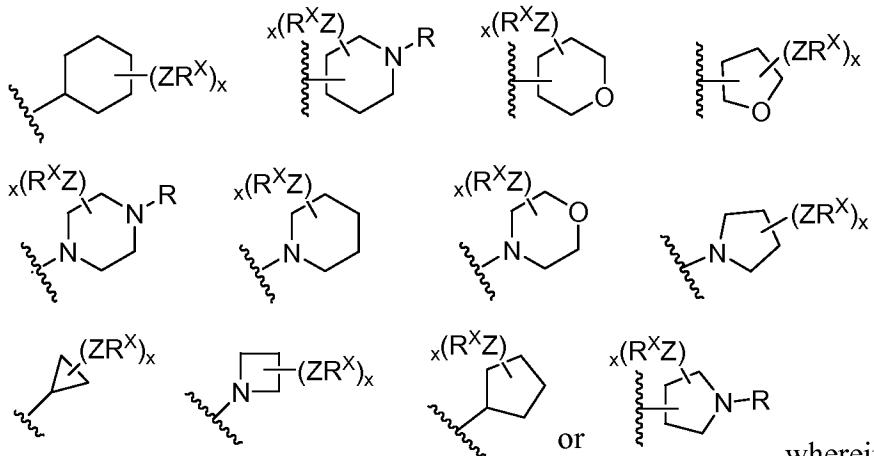
Ar¹ is selected from



Applicants: Jeremy Green et al.
 Application No.: 10/808,678



Cy¹ is selected from



wherein;

Ar¹ and Cy¹ are each optionally substituted with up to 5 occurrences of Z-R^X; wherein each occurrence of Z is independently a bond or a C₁₋₆ alkylidene chain, wherein up to two non-adjacent methylene units of Z are optionally replaced by -S-, -O-, -N(R)-, -N(R)C(O)-, -C(O)N(R)-, -SO₂N(R)-, or -N(R)SO₂-;

each occurrence of R^X is independently selected from -R', halogen, NO₂, CN, -OR', -SR', or -N(R')₂,

each occurrence of R is independently hydrogen or a C₁₋₆ aliphatic group; and

each occurrence of R' is independently hydrogen, a C₁₋₆ aliphatic group, a C₆₋₁₀ aryl ring, a heteroaryl ring having 5-10 ring atoms, or a heterocyclyl ring having 3-10 ring atoms; or

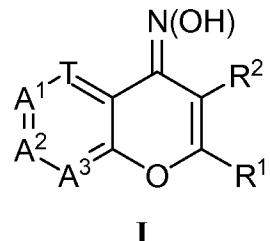
R and R' or two occurrences of either R or R' are taken together with the atoms to which they are bound to form an optionally substituted 5-8 membered saturated, partially unsaturated, or aryl ring having 0-4 heteroatoms independently selected from nitrogen, oxygen, or sulfur; or

Applicants: Jeremy Green et al.
Application No.: 10/808,678

two occurrences of either R' or R on the same nitrogen are taken together with the nitrogen atom to which they are bound to form an optionally substituted 5-8 membered saturated, partially unsaturated, or aryl ring having 1-4 heteroatoms independently selected from nitrogen, oxygen, or sulfur.

48-49. (Canceled)

50. (Currently amended) A method of inhibiting c-MET kinase activity in a biological sample, wherein said biological sample is selected from a cell culture, biopsied material obtained from a mammal, saliva, urine, feces, semen, or tears, or an extract thereof; which method comprises contacting said biological sample with a composition according to claim 47 or a compound of formula I:



or a pharmaceutically acceptable salt thereof, wherein:

R¹ is Ar¹;

R² is hydrogen;

T is CH;

A¹ is C-halogen, C-CN, or C-R;

each of A² and A³ is, independently, CR⁴;

R⁴ is selected from halogen, NO₂, CN, -(L)_mR, -(L)_mAr¹, or -(L)_mCy¹; or

two R⁴ groups on adjacent atoms are taken together to form an optionally substituted 5-7 membered partially unsaturated or fully unsaturated ring having 0-3 heteroatoms independently selected from oxygen, sulfur, or nitrogen, wherein each ring formed by

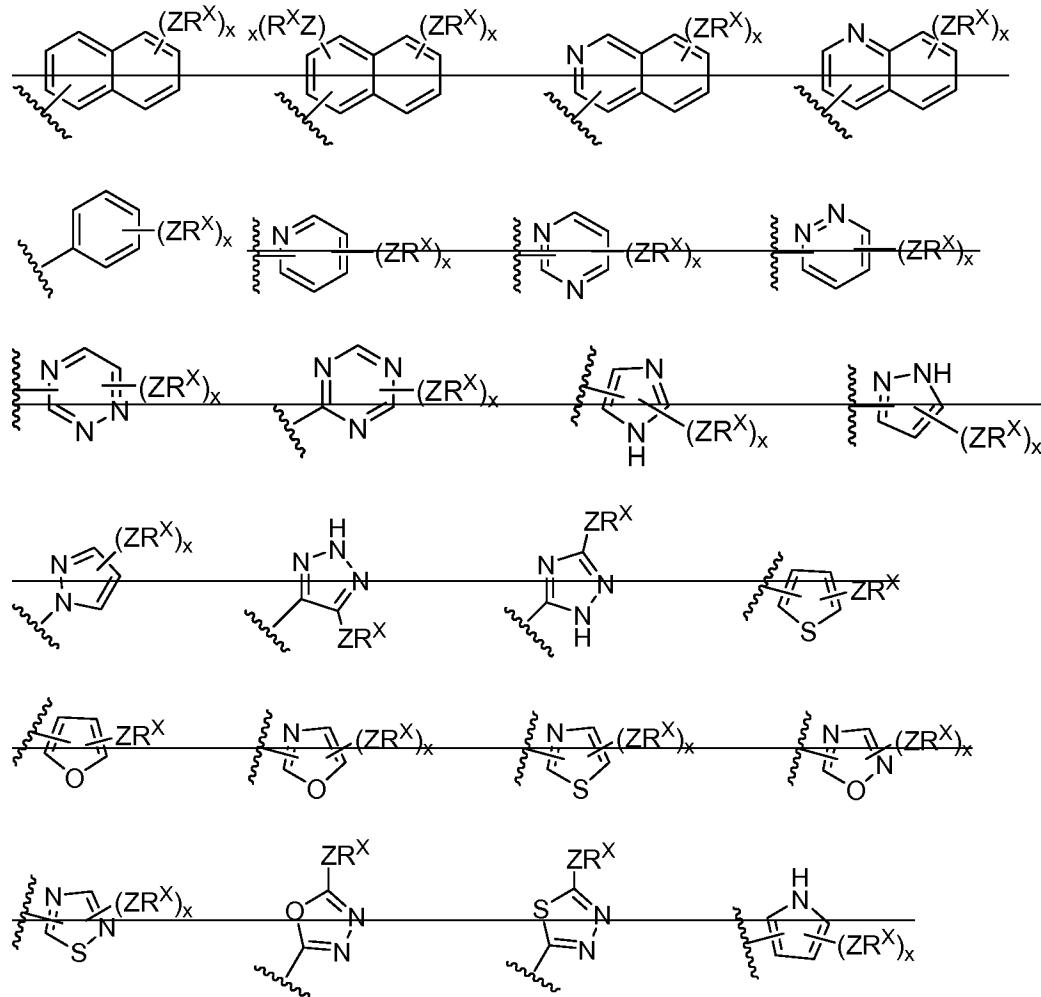
Applicants: Jeremy Green et al.
Application No.: 10/808,678

two R⁴ groups on adjacent atoms taken together is optionally substituted with up to 4 occurrences of Z-R^X;

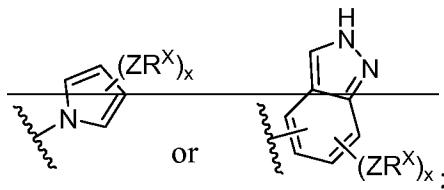
L is a C₁₋₆ alkylidene chain wherein one methylene unit of L is optionally replaced by -O-, -N(R)-, -N(R)C(O)-, -C(O)-, -C(O)N(R)-, -SO₂N(R)-, or -N(R)SO₂-;

m is 0 or 1;

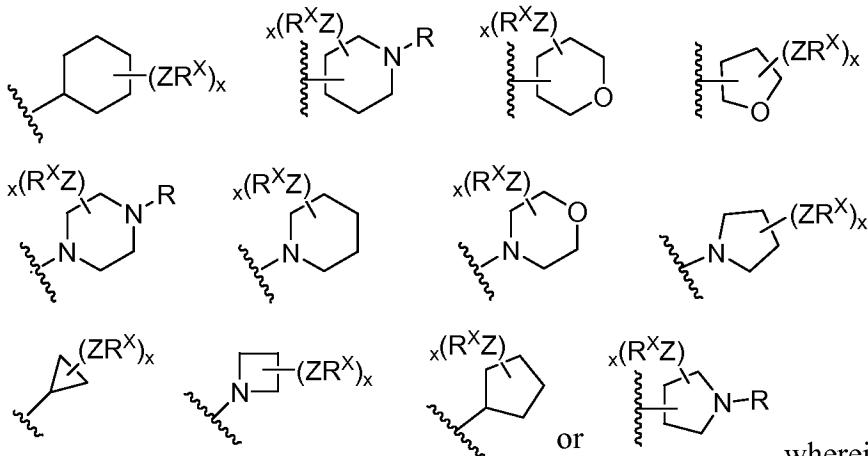
Ar¹ is selected from



Applicants: Jeremy Green et al.
 Application No.: 10/808,678



Cy¹ is selected from



wherein;

Ar¹ and Cy¹ are each optionally substituted with up to 5 occurrences of Z-R^X; wherein each occurrence of Z is independently a bond or a C₁₋₆ alkylidene chain, wherein up to two non-adjacent methylene units of Z are optionally replaced by -S-, -O-, -N(R)-, -N(R)C(O)-, -C(O)N(R)-, -SO₂N(R)-, or -N(R)SO₂-;

each occurrence of R^X is independently selected from -R', halogen, NO₂, CN, -OR', -SR', or -N(R')₂,

each occurrence of R is independently hydrogen or a C₁₋₆ aliphatic group; and

each occurrence of R' is independently hydrogen, a C₁₋₆ aliphatic group, a C₆₋₁₀ aryl ring, a heteroaryl ring having 5-10 ring atoms, or a heterocyclyl ring having 3-10 ring atoms; or

R and R' or two occurrences of either R or R' are taken together with the atoms to which they are bound to form an optionally substituted 5-8 membered saturated, partially unsaturated, or aryl ring having 0-4 heteroatoms independently selected from nitrogen, oxygen, or sulfur; or

Applicants: Jeremy Green et al.
Application No.: 10/808,678

two occurrences of either R' or R on the same nitrogen are taken together with the nitrogen atom to which they are bound to form an optionally substituted 5-8 membered saturated, partially unsaturated, or aryl ring having 1-4 heteroatoms independently selected from nitrogen, oxygen, or sulfur.

51. (Canceled)

52. (Withdrawn) A method of treating or lessening the severity of a disease or condition in a patient selected from gastric cancer, pancreatic cancer, ovarian cancer, breast cancer, or prostate cancer comprising the step of administering to said patient a composition of claim 47.

53. (Withdrawn) The method according to claim 52, comprising the additional step of administering to said patient an additional therapeutic agent selected from mechlorethamine, chlorambucil, cyclophosphamide, melphalan, ifosfamide, methotrexate, 6-mercaptopurine, 5-fluorouracil, cytarabine, gemcitabine, vinblastine, vincristine, vinorelbine, paclitaxel, etoposide, irinotecan, topotecan, doxorubicin, bleomycin, mitomycin, carmustine, lomustine, cisplatin, carboplatin, asparaginase, and tamoxifen, leuprolide, flutamide, megestrol, imatinib (GleevecTM), adriamycin, dexamethasone, or cyclophosphamide, wherein:

said additional therapeutic agent is appropriate for the disease being treated; and
said additional therapeutic agent is administered together with said composition
as a single dosage form or separately from said composition as part of a
multiple dosage form.

54-80. (Canceled)

Applicants: Jeremy Green et al.
Application No.: 10/808,678

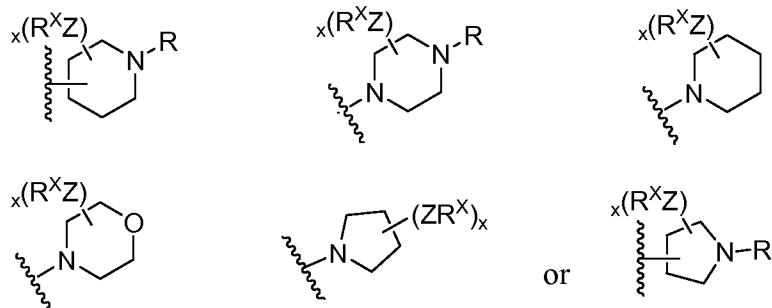
81. (Previously presented) The composition according to claim 47, wherein A² is CR⁴ and R⁴ is halogen, CN, -(L)_mR, -(L)_mAr¹, or -(L)_mCy¹.

82. (Canceled)

83. (Previously presented) The composition according to claim 81, wherein A² is CR⁴ and R⁴ is halogen or R.

84. (Previously presented) The composition according to claim 81, wherein A² is CR⁴ and R⁴ is -(L)_mR, wherein L is -O- or -N(R)-.

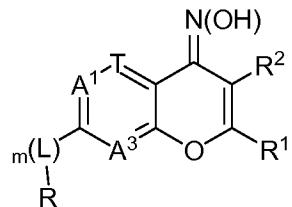
85. (Previously presented) The composition according to claim 81, wherein A² is CR⁴, R⁴ is -(L)_mCy¹, m is 0 and Cy¹ is



86. (Canceled)

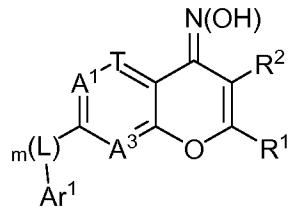
Applicants: Jeremy Green et al.
Application No.: 10/808,678

87. (Previously presented) The composition according to claim 81, wherein A² is CR⁴, R⁴ is -(L)_mR, and compounds have the formula **IE-1**:



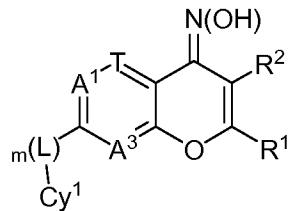
IE-1.

88. (Previously presented) The composition according to claim 81, wherein A² is CR⁴, R⁴ is -(L)_mAr¹, and compounds have the formula **IE-2**:



IE-2 .

89. (Previously presented) The composition according to claim 81, wherein A² is CR⁴, R⁴ is -(L)_mCy¹, and compounds have the formula **IE-3**:



IE-3 .

Applicants: Jeremy Green et al.
Application No.: 10/808,678

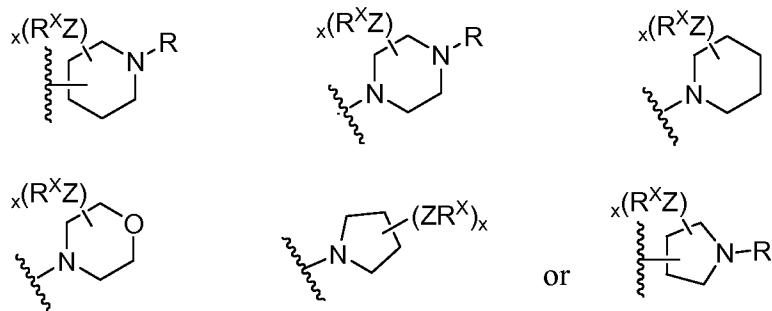
90. (Previously presented) The composition according to claim 47, wherein A³ is CR⁴ and R⁴ is halogen, CN, -(L)_mR, -(L)_mAr¹, or -(L)_mCy¹.

91. (Canceled)

92. (Previously presented) The composition according to claim 90, wherein A³ is CR⁴ and R⁴ is halogen or R.

93. (Previously presented) The composition according to claim 90, wherein A³ is CR⁴ and R⁴ is -(L)_mR, wherein L is -O- or -N(R)-.

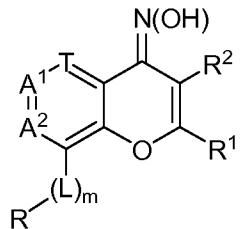
94. (Previously presented) The composition according to claim 90, A³ is CR⁴, R⁴ is -(L)_mCy¹, m is 0 and Cy¹ is



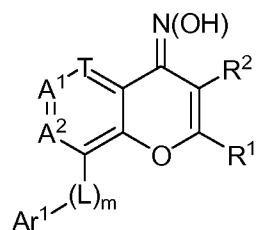
95. (Canceled)

Applicants: Jeremy Green et al.
Application No.: 10/808,678

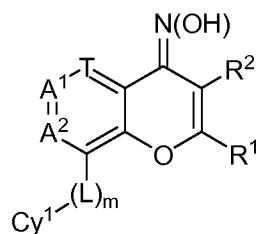
96. (Previously presented) The composition according to claim 90, wherein A³ is CR⁴, R⁴ is -(L)_mR, and compounds have the formula **IF-1**:



97. (Previously presented) The composition according to claim 90, wherein A³ is CR⁴, R⁴ is -(L)_mAr¹, and compounds have the formula **IF-2**:



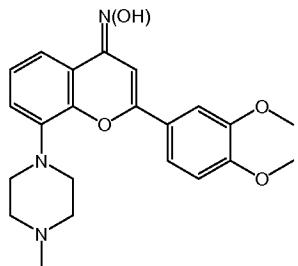
98. (Previously presented) The composition according to claim 90, wherein A³ is CR⁴, R⁴ is -(L)_mCy¹, and compounds have the formula **IF-3**:



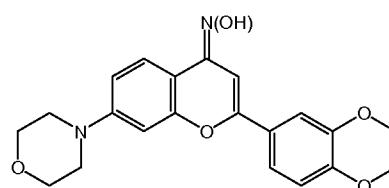
Applicants: Jeremy Green et al.
Application No.: 10/808,678

99-100. (Canceled)

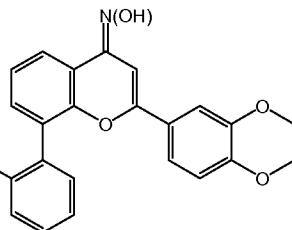
101. (Currently amended) The composition according to claim 47, selected from one of the following compounds:



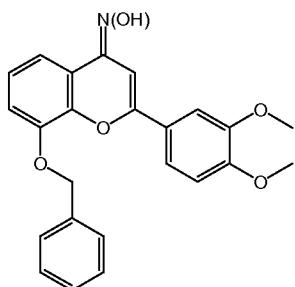
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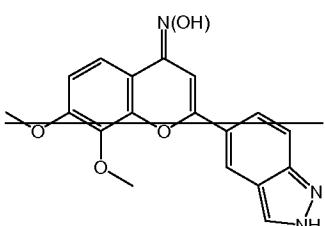
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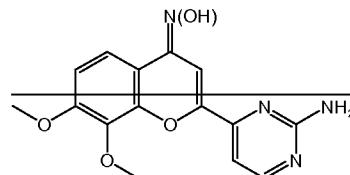
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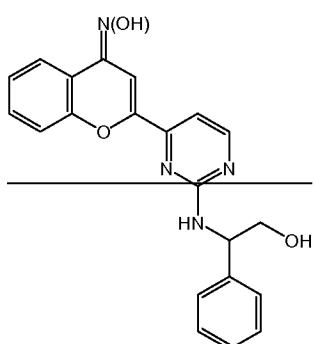
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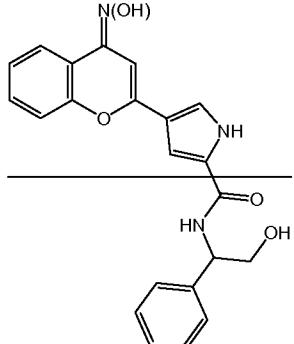
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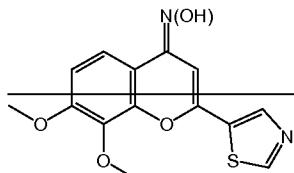
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[[I-7]],

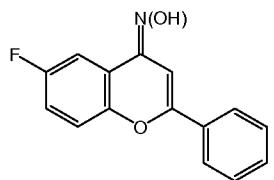


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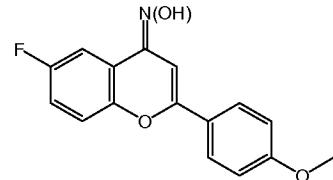


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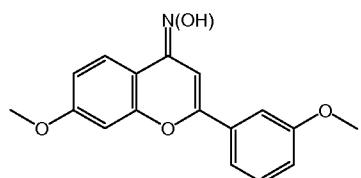
Applicants: Jeremy Green et al.
Application No.: 10/808,678



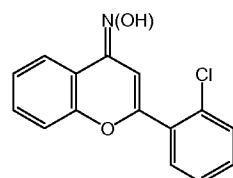
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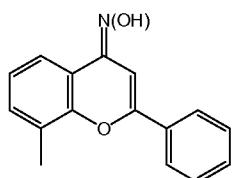
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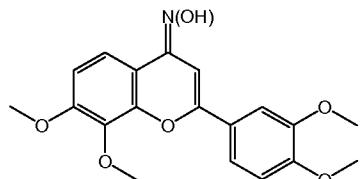
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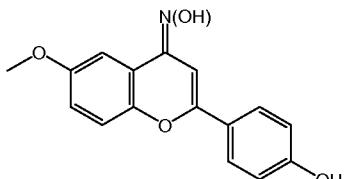
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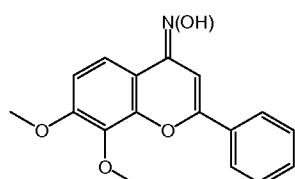
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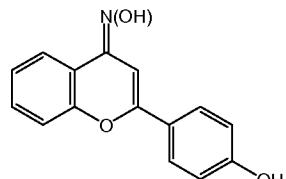
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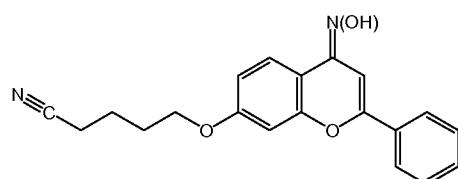
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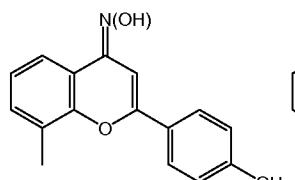
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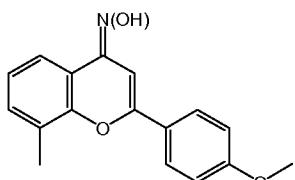
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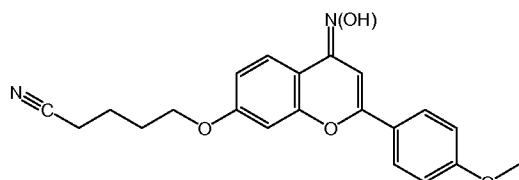
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I-22,

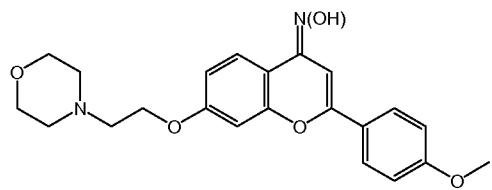


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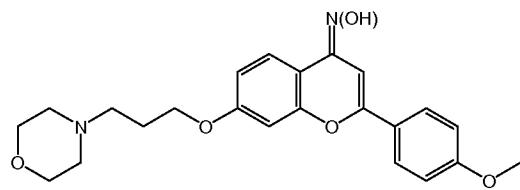


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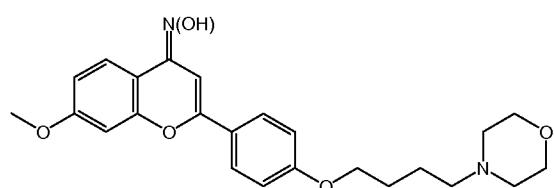
Applicants: Jeremy Green et al.
Application No.: 10/808,678



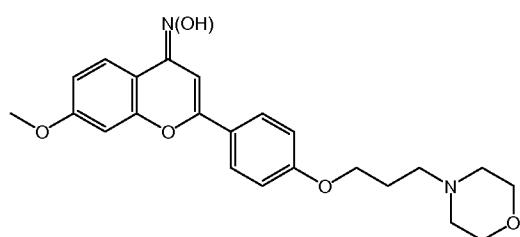
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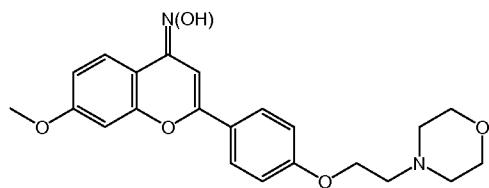
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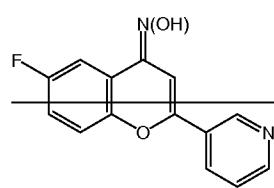
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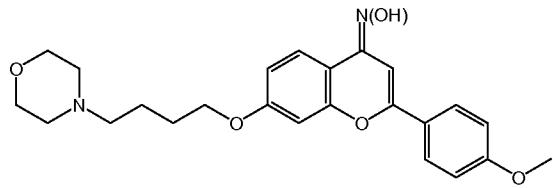
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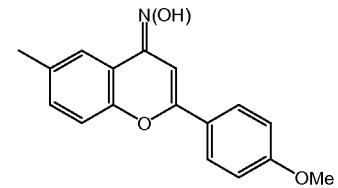
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I-30,

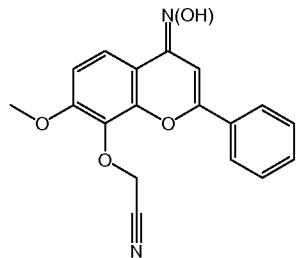


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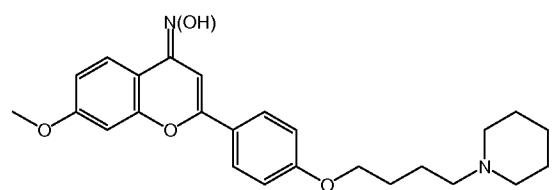


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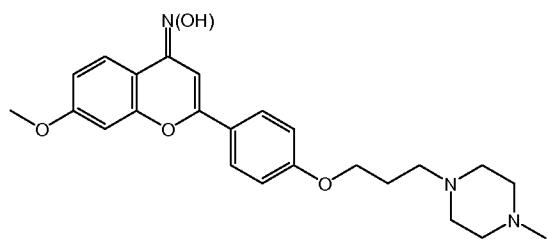
Applicants: Jeremy Green et al.
Application No.: 10/808,678



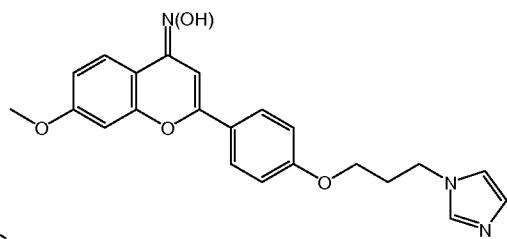
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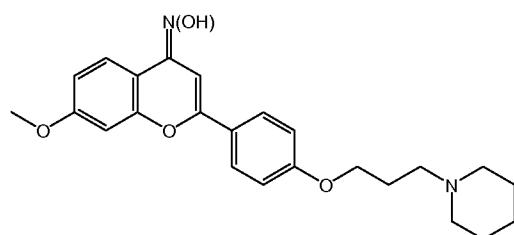
I-34,



I-35,



I-36,



or I-37 .